

UNITED STATE DEPARTMENT OF COMMERCE **United States Patent and Trademark Office**

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	APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR		ATTORNEY DOCKET NO.	
	09/757,2	12 01/09	/01 SEDRANI		R	100-8024C/C:
Г	0010 9 5	001095 · HM12/08			EXAMINER	
	THOMAS HOXIE NOVARTIS CORPORATION				CEPERI ART UNIT	EY.M PAPER NUMBER
	564 MORR	ND TRADEMAN IS AVENUE J 07901-10:			1641 DATE MAILED:	3
						08/07/01

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trademarks

اور بتهتریاندا م		Application No.	Applicant(s)					
,,	. •	09/757,212	SEDRANI ET AL.					
	Office Action Summary	Examiner	Art Unit					
		Mary E. (Molly) Ceperley	1641					
	The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply							
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). - Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b). Status								
1) 🗌	Responsive to communication(s) filed on							
2a) <u></u>	This action is FINAL . 2b)⊠ Th	is action is non-final.						
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.								
Dispositi	Disposition of Claims							
4)⊠	4)⊠ Claim(s) <u>1-14</u> is/are pending in the application.							
4a) Of the above claim(s) is/are withdrawn from consideration.								
5) Claim(s) is/are allowed.								
6)	6) ☐ Claim(s) is/are rejected.							
7) 🗌	7) Claim(s) is/are objected to.							
8)⊠	8) Claim(s) 1-14 are subject to restriction and/or election requirement.							
Application Papers								
9)☐ The specification is objected to by the Examiner.								
10) ☐ The drawing(s) filed on is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.								
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).								
11) The proposed drawing correction filed on is: a) approved b) disapproved by the Examiner.								
If approved, corrected drawings are required in reply to this Office action.								
12)☐ The oath or declaration is objected to by the Examiner.								
Priority u	nder 35 U.S.C. §§ 119 and 120	•						
13)🛛	13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).							
a) ☐ All b) ☐ Some * c) ☐ None of:								
	1. Certified copies of the priority documents have been received.							
2. Certified copies of the priority documents have been received in Application No								
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 								
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).								
a) The translation of the foreign language provisional application has been received.								
15)⊠ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.								
Attachment(s)								
2) Notice	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449) Paper No(s) _	5) Notice of Informal	y (PTO-413) Paper No(s) Patent Application (PTO-152)					
J.S. Patent and Tr PTO-326 (Re		ction Summary	Part of Paper No. 3					

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المراجعة المراجعة

1. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in response to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-7, 13 and 14, drawn to monoclonal antibodies capable of specifically recognizing a rapamycin, a hybridoma cell line capable of producing said monoclonal antibody and a kit comprising said monoclonal antibody.

Group II, claim(s) 8-10, drawn to immunogenic conjugates comprising a rapamycin portion and a protein portion.

Group III, claims 11 and 12, drawn to a rapamycin having an activated coupling group.

The inventions listed as Groups I-III do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

Groups I-III each possess a special technical feature which is structurally, biologically and chemically different and therefore patentably distinct from the special technical features of the other Groups. The special technical feature of Group I is an antibody which specifically recognizes rapamycin. The special technical feature of Group II is an immunogenic conjugate which does not use the antibody of Group I, i.e. it comprises different components, a rapamycin

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portion and a protein portion. The special technical feature of Group III is a rapamycin having an activated coupling group. This functionalized rapamycin is a different product from that of Groups I and II. Therefore, Groups I-III do not relate to a single general inventive concept i.e. drug-protein conjugates, antibodies, and activated drug molecules of the instant application are structurally, biologically and chemically distinct.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

2. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Mary E. (Molly) Ceperley whose telephone number is (703) 308-4239. The examiner can normally be reached from 8 a.m. to 5 p.m.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le, can be reached on (703) 305-3399. The fax phone number for the organization where this application or proceeding is assigned is (703) 305-7230.

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Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-1234.

August 7, 2001

Mary E. Ceperley

Mary E. Ceperley

Primary Examiner

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